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Declarations under Rule 4.17:

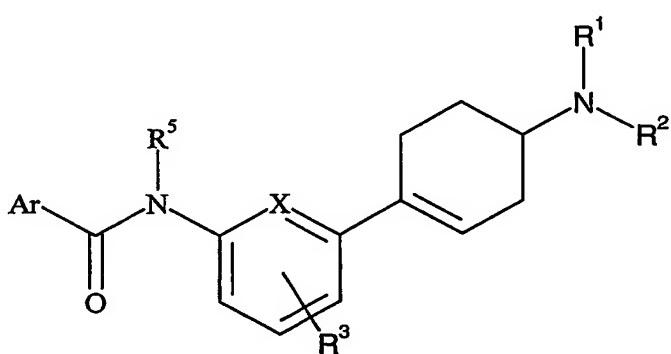
— as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, ARIPO patent (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)

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(54) Title: SUBSTITUTED (4-AMINOCYCLOHEXEN-1-YL)PHENYL AND (4-AMINOCYCLOHEXEN-1-YL)PYRIDINYL COMPOUNDS AS 5-HT_{1F} AGONISTS

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(57) Abstract: The present invention relates to compounds of formula (I) or a pharmaceutically acceptable acid addition salt thereof, where; X is -C(R⁴)= or -N=; Ar is phenyl, substituted phenyl, heterocycle, or substituted heterocycle; R¹ and R² are independently hydrogen or C₁-C₃ alkyl; R³ is hydrogen, fluoro, or methyl; when X is -C(R⁴)=, R⁴ is hydrogen, fluoro, or methyl, provided that no more than one of R³ and R⁴ may be other than hydrogen; and R⁵ is hydrogen, methyl, or ethyl. The compounds of the present invention are useful for activating 5-HT_{1F} receptors, inhibiting dural

protein extravasation, and for the treatment or prevention of migraine in a mammal.